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- (71) Applicant (for all designated States except US): **3M INNOVATIVE PROPERTIES COMPANY** [US/US]; 3M Center, Post Office Box 33427, Saint Paul, MN 55133-3427 (US).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): **SLOWEY, Alexander D.**, [GB/GB]; 61 Kennedy Drive, Stapleford, Nottinghamshire NG9 8HW (GB). **BOSWELL, Susannah C.**, [GB/GB]; 62 Lindisfarne Way, East Hunsbury, Northamptonshire NN4 0WJ (GB). **JINKS, Philip A.**, [GB/GB]; 41 Highfields Drive, Loughborough, Leicestershire LE11 3JS (GB).
- (74) Agents: **RINGSRED, Ted K.**, et al.; Office of Intellectual Property Counsel, Post Office Box 33427, Saint Paul, MN 55133-3427 (US).
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(54) Title: **FORMOTEROI. AND MOMETASONE AEROSOL FORMULATIONS**

(57) Abstract: A pharmaceutical aerosol formulation comprising particles of (a) formoterol or a pharmaceutically acceptable salt, solvate or physiologically functional derivative thereof and (b) mometasone or a pharmaceutically acceptable salt, solvate, or physiologically functional derivative thereof dispersed in a propellant selected from the group consisting of 1,1,1,2-tetrafluoroethane, 1,1,1,2,3,3,3-heptafluoropropane and a mixture thereof, and a bulking agent having a mass median diameter of less than one micron.

FORMOTEROL AND MOMETASONE AEROSOL FORMULATIONS

5 Field of the Invention

This invention relates to medicinal aerosol formulations and in particular to suspension aerosol formulations containing formoterol and mometasone and a nano-particulate auxiliary powder suitable for administration to the respiratory tract.

Background

Formoterol, N-[2-hydroxy-5-(1-hydroxy-2-(2-(4-methoxyphenyl)-1-methylethyl) amino)ethyl) phenyl] formamide, particularly in the form of its fumarate salt, is a bronchodilator used in the treatment of inflammatory or obstructive airways diseases.

EP 057 401 and US 4,472,393 describe mometasone i. e. 9,21-dichloro-11,17-dihydroxy-16 α -methylpregna-1, 4-diene-3, 20-dione, esters thereof such as mometasone furoate i. e. (11 β , 16 α)-9, 21-dichloro-17- [(2-furanylcarbonyl) oxy]-11- hydroxy-16-methylpregna-1, 4-diene-3,20-dione, and pharmaceutical formulations thereof. Mometasone is an antiinflammatory corticosteroid, which is now used clinically in the treatment of respiratory disorders.

WO 00/51591 describes medicaments and pharmaceutical compositions containing, separately or together, (A) formoterol or a pharmaceutically acceptable salt thereof or a solvate of formoterol or a solvate of said salt and (B) mometasone furoate, for simultaneous, sequential or separate administration in the treatment of an inflammatory or obstructive airways disease. As an example for a suspension aerosol formulation for a metered dose inhaler, WO 00/51591

discloses a composition consisting of 0.012 % by weight formoterol fumarate dihydrate, 0.250 % by weight mometasone furoate, 2.5% by weight ethanol, 60.768 % by weight 1,1,1,2,3,3,3-heptafluoropropane (HFC 227) and 36.740 % by weight 1,1,1,2-tetrafluoroethane (HFC 134a).

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WO 00/53187 discloses suspension aerosol formulations consisting of micronized formoterol fumarate dihydrate (6 μg), micronized mometasone furoate monohydrate (100 μg), oleic acid (0.005 % based on propellant), ethanol (2% based on propellant) and HFC 134a (up to 25, 50 or 100 μl) as well as suspension formulations consisting of micronized formoterol fumarate dihydrate (12 μg), micronized mometasone furoate monohydrate (200 μg), oleic acid (0.01 % based on propellant), ethanol (3% based on propellant) and HFC 227/HFC 134a (15/85, up to 25, 50 or 100 μl).

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WO 01/78744 discloses suspension aerosol formulations consisting of micronized formoterol fumarate or the dihydrate thereof (6 to 24 μg per actuation), micronized mometasone furoate (100 or 200 μg per actuation) and HFC 134a.

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An important aspect in formulating pharmaceutical compositions which have a micronized drug dispersed in a propellant is the homogeneity of the dispersed drug. The drug can sediment or cream depending on the density difference between drug and propellant, or it can flocculate, which requires some degree of agitation to deflocculate it. Inhomogeneity of the dispersed drug may in turn lead to delivery of incorrect doses when the formulation is dispensed from the metering valve. This aspect represents a challenge when formulating suspensions of a single drug, but is particularly challenging for a combination of drugs, like formoterol and mometasone, in which each drug has a significantly different potency and thus concentration in formulation. Furthermore, the densities of the two drugs differ appreciably, e.g. formoterol fumarate has a density of 1.3 g cm^{-3} , while mometasone furoate has a density of 1.4 g cm^{-3} .

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Summary of the Invention

We have found that the use of a particulate bulking agent having a mass median diameter of less than one micron in aerosol compositions comprising formoterol and mometasone dispersed in HFC 134a, HFC 227 or mixtures thereof, allows the provision of dispersions of both drugs showing desirably high physical stability and homogeneity.

The present invention provides a pharmaceutical aerosol formulation comprising particles of (a) formoterol or a pharmaceutically acceptable salt, solvate or physiologically functional derivative thereof and (b) mometasone or a pharmaceutically acceptable salt, solvate, or physiologically functional derivative thereof dispersed in propellant HFC 134a and/or HFC 227 and a bulking agent having a mass median diameter of less than one micron.

In particular, it has been found that the application of such a nano-sized bulking agent having a mass median diameter of less than one micron aids in minimizing the tendency of both formoterol and mometasone to cream or sediment, independent of the density difference of the respective drug to one another and to the propellant. More particularly, it has been found that the application of such a nano-sized bulking agent aids in maintaining a high sediment volume (i.e. minimizing a dense packing of the sediment) and/or the formation of a voluminous, loosely flocculated matrix, enhancing the re-dispersion and/or deflocculation of the two drugs upon agitation.

The use of a particulate bulking agent having a mass median diameter of less than one micron in pharmaceutical aerosol formulations comprising a suspension of drug particles in a propellant is disclosed in commonly assigned co-pending application PCT US01/30575.

Formulations according to the invention are particularly suitable for use in inhalation therapy, in which a therapeutically effective amount of the formulation is

delivered to the lung by oral or nasal inhalation, more particularly for prophylaxis or treatment of a clinical condition for which a selective β_2 -adrenoreceptor agonist and/or antiinflammatory corticosteroid is indicated. The present invention also provides a method for the prophylaxis or treatment of a clinical condition in a mammal, such as a human, for which a selective β_2 -adrenoreceptor agonist and/or antiinflammatory corticosteroid is indicated, which comprises administration via inhalation a therapeutically effective amount of the formulation as described above. In particular, the present invention provides such methods for the prophylaxis or treatment of a disease associated with reversible airways obstruction such as asthma, chronic obstructive pulmonary disease (COPD), respiratory tract infection or upper respiratory tract disease.

In another aspect of the present invention, there is provided a dispenser comprising an aerosol vial equipped with a dispensing valve containing a formulation as described above.

Detailed Description

It is to be understood that the present invention covers all combinations of particular and preferred aspects of the invention described herein.

As would be appreciated by the skilled person, formoterol includes two asymmetric centres, and mometasone contains several asymmetric centres. The present invention includes each isomer of formoterol either in substantially pure form or admixed in any proportions or a racemic mixture, particularly the (R, R)-isomer as well as each isomer of mometasone either in substantially pure form or admixed in any proportions. The enantiomers of formoterol have been described previously, for example, in WO 98/21175 and US5795564.

By the term "physiologically functional derivative" is meant a chemical derivative of formoterol or mometasone having the same physiological function as the free compound, for example, by being convertible in the body thereto. According to the

present invention, examples of physiologically functional derivatives include esters.

Suitable salts according to the invention include those formed with both organic and inorganic acids. Pharmaceutical acceptable acid addition salts include but are not limited to those formed from hydrochloric, hydrobromic, sulphuric, citric, tartaric, phosphoric, lactic, pyruvic, acetic, trifluoroacetic, succinic, oxalic, fumaric, maleic, oxaloacetic, methanesulphonic, ethanesulphonic, p-toluenesulphonic, benzenesulphonic, isethionic, and naphthalenecarboxylic, such as 1-hydroxy-2-naphthalenecarboxylic acids.

Pharmaceutically acceptable esters of formoterol or mometasone may have a hydroxyl group converted to a C₁₋₆ alkyl, aryl, aryl C₁₋₆ alkyl, hetaryl (such as furanyl) or amino acid ester.

In preferred embodiments of the invention, formoterol fumarate (suitably as in the form of the dihydrate) is applied in combination with mometasone furoate (suitably as in the form of the monohydrate or more particularly in the anhydrous form).

Hereinafter, the term "formoterol" is understood to include formoterol or a pharmaceutical acceptable salt, solvate, or physiologically functional derivative thereof, preferably formoterol fumarate, while the term "mometasone" includes mometasone, or a pharmaceutical acceptable salt, solvate, or physiologically functional derivative thereof, preferably mometasone furoate, unless specified otherwise.

Formoterol and mometasone are generally present in a formulation of the invention in a therapeutically effective amount. The amount of formoterol and mometasone which is required to achieve a therapeutic effect will, of course, vary with the particular compound, the subject under treatment, and the particular disorder or disease being treated. Suitably, the pharmaceutical formulations which are suitable for inhalation according to the invention comprise formoterol

and mometasone in amounts such that one or two actuations provide a therapeutically effective dose, for example, a dose of formoterol of 1 mcg to 50 mcg, preferably 3 mcg to 25 mcg, more preferably 4 mcg to 12 mcg and a dose of mometasone of 50 mcg to 1.6mg, preferably 50 mcg to 1 mg, more preferably, 50 mcg to 200 mcg. Various dosing of the individual drugs can be advantageously combined for particular disorders or subjects under treatment. For example a formoterol dosing, such as 6 mcg per actuation can be combined with a low strength dosage of mometasone, e.g. 50 mcg per actuation, or a higher strength dosage, e.g. 200 mcg per actuation.

Preferably formoterol constitutes about 0.06 to about 0.60 mg per ml, more preferably about 0.08 to about 0.30 mg per ml, even more preferably about 0.1 to about 0.2 mg per ml, most preferably about 0.132 mg per ml of the formulation.

Mometasone preferably constitutes about 0.5 to about 15 mg per ml of the formulation. For high strength formulations mometasone constitutes preferably about 3 to about 15 mg per ml, more preferably about 4 to about 10 mg per ml, most preferably about 4.0 to 6.0 mg per ml of the formulation. For aerosol formulations including lower strength doses of mometasone, mometasone preferably constitutes about 0.5 to about 3.0 mg per ml, more preferably about 0.8 to about 2.5 mg per ml, most preferably about 1.00 to 1.25 mg per ml of the formulation.

The particles of formoterol or mometasone are generally micronised particles or particles processed by other methods, preferably having a mass median diameter equal to or greater than 1 micron, more particularly from 1 to 10 micron, even more particularly from 1 to 5 micron. Smaller particles having a mass median diameter of less than one micron may also be suitable.

The bulking agent having a mass median diameter of less than one micron, improves the stability of the suspension of formoterol and mometasone particles.

It is not necessary for the surface of the bulking agent or the drug to be coated with a surface modifier to achieve improved stability.

The mass median diameter of the bulking agent can advantageously be as low as 300 nanometers, more desirably as low as 250 nanometers, even more desirably the mass median diameter is in the range of 100 to 250 nanometers and most desirably in the range of 150 to 200 nanometers.

Mass median diameter (which is equivalent to volume median diameter) can be determined using any conventional particle size measurement method known to those skilled in the art. Suitable methods include for example laser diffraction, photon correlation spectroscopy (e.g. using a spectrometer available under the trade designation Brookhaven PCS from Brookhaven Inc.), spinning disc centrifuging (using an instrument available under the trade designation CPS Disc Centrifuge from Chemical Process Specialists Inc.) and scanning electron microscopy (SEM). Mass median diameter is preferably determined by laser diffraction, photon correlation spectroscopy or spinning disc centrifuging, more preferably by laser diffraction, more particularly laser diffraction using an analyser available under the trade designation Malvern Mastersizer 2000 laser light diffraction particle size analyser from Malvern Instruments Ltd.

Preferred bulking agents include lactose, DL-alanine, ascorbic acid, glucose, sucrose D(+)-trehalose as well as their various hydrates, anomers and/or enantiomers. Lactose including its various forms, such as α -lactose monohydrate and β -lactose and alanine are more preferred. Lactose, in particular in its α -lactose monohydrate form, is most preferred as a bulking agent due to e.g. processing considerations. Other suitable bulking agents include other saccharides e.g. D-galactose, maltose, D(+)-raffinose pentahydrate, sodium saccharin, polysaccharides e.g. starches, modified celluloses, dextrans or dextrins, other amino acids e.g. glycine, salts e.g. sodium chloride, calcium carbonate, sodium tartrate, calcium lactate, or other organic compounds e.g. urea or propylidone.

Based on the amount of formoterol in the formulation, the weight ratio of formoterol to bulking agent is generally in the range 1:0.1 to 1:25, preferably 1:2 to 1:15, even more preferably 1:4 to 1:12 and most preferably 1:8 to 1:10.

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For formulations including a low strength dosage of mometasone, e.g. 50 mcg per actuation, the weight ratio of total drug (formoterol and mometasone) to bulking agent is generally in the range 1:0.01 to 1:3, preferably 1:0.2 to 1:1.8, even more preferably 1:0.4 to 1:4 and most preferably about 1: 0.94 to 1:1.17.

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For formulations including a high strength dosage of mometasone, e.g. 200 mcg per actuation, the weight ratio of total drug (formoterol and mometasone) to bulking agent is generally in the range 1:0.002 to 1:6, preferably 1:0.05 to 1:0.4, even more preferably 1:0.1 to 1:3 and most preferably about 1:0.20 to 1:0.25.

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To further enhance wetting characteristics of the formulation, and thus further minimizing the deposition of drug onto the metal parts of the container closure system and onto stainless steel manufacturing equipment during manufacture, the aerosol formulations of the invention may desirably contain ethanol. When ethanol is included in a formulation, it is generally present in an amount in the range 0.1 to 5% by weight of the total formulation, preferably from about 0.5 to about 4% by weight, more preferably from about 1 to about 3% by weight, most preferably about 2 % by weight.

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The aerosol formulations of the invention may preferably contain surfactant for e.g. imparting a flocculant effect which may allow less migration of the drugs to and from the metering chamber. When surfactant is included in a formulation it is generally present in an amount of about 0.001 % to 0.050 % by weight of the formulation. Suitable surfactants are well known in the art and include sorbitan trioleate, oleic acid and lecithin. Surfactants, such as oligolactic acid derivatives disclosed in WO94/21228 and WO94/21229, and other surfactants disclosed in the literature may be used. As a surfactant oleic acid is preferred.

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The bulking agent may be reduced to the required particle size by any convenient method, e.g. grinding, air-jet milling etc. Preferably the bulking agent is reduced to nanoparticle size in a high pressure homogenizer, such as the commercially available Avestin Emulsiflex homogenizers and the Microfluidics Microfluidizer homogenizers. In the processing with high pressure homogenizers, certain bulking agents can be reduced to the desired particle size using lower pressures than that applied for other bulking agents. For example, it has been found that lactose, more specifically α -lactose monohydrate, can be effectively reduced to the desired particle size using pressures between about 10,000 and about 21,000 psi, while for effective particle size reduction of alanine or sucrose higher pressures of about 25,000 psi for repeated passes were applied.

The bulking agent may be prepared in a slurring aid which is a low volatility solvent such as ethanol. It may be prepared in a slurring aid which is a component of the final aerosol formulation, or it may be prepared in a solvent that is subsequently removed or exchanged with a component of the formulation by some process such as centrifugation and decanting, dialysis, evaporation etc.

It is particularly convenient to use a slurring aid in the high pressure homogenizer which is a low volatility component of the aerosol formulation and after particle size reduction has been achieved the slurry may be adjusted if necessary, e.g. concentrated by centrifugation, decanting etc. Whilst it has been found that slurries with excessively high powder loadings may be difficult to process due to their rheological properties, it is generally advantageous to process slurries with powder loading concentrations which approach this processing limit in order to achieve the desired particle size distribution in the shortest processing time. Thus, the weight ratio of liquid:solid is generally in the range 5:1 to 100:1, preferably 5:1 to 20:1, and most preferably 8:1 to 10:1.

The present invention also provides a method of preparing a formulation according to the invention, the method comprising the steps of (i) forming a slurry of bulking agent with a component of the formulation; (ii) subjecting the slurry to

high pressure homogenization; and (iii) combining the resulting slurry with other components of the aerosol formulation.

For formulations containing ethanol, the slurry of bulking agent may be
5 advantageously prepared with an appropriate amount of ethanol. The slurry is
subjected to high pressure homogenization prior to adding it to the remainder of
the formulation. During manufacture, formoterol and mometasone are typically,
initially dispersed with an appropriate amount of HFC 134a and/or HFC 227 and,
if applicable or desired ethanol, i.e. such an amount of ethanol to aid dispersion
0 but not so as to result in excessive partial solubilisation of either of the drugs, to
prepare a cold concentrate. This dispersion, typically after high shear mixing, is
then added to a second appropriate amount of HFC 134a and/or HFC 227 in
liquid form, chilled to below its boiling point or range. Thereafter the
(homogenized) bulking agent slurry is added. Alternatively the bulking agent slurry
15 may be added to the second appropriate amount of propellant before the drug
dispersion is added. If surfactant is used, it is preferably dissolved at ambient
temperature in an appropriate amount of ethanol, and it is either added to the
drug containing dispersion or the bulking agent slurry.

20 Alternatively, the formulations made by prepared by single stage or two stage
pressure filling.

Dispensers comprising an aerosol vial equipped with conventional dispensing
valves, preferably metered dose valves, can be used to deliver formulations of the
25 invention. Conventional dispensers and aerosol vials can be used to contain a
formulation of the invention. However it has been found that certain vials enhance
the chemical stability of certain formulations of the invention. Therefore it is
preferred to contain a formulation of the invention within a glass aerosol vial or a
metal, in particular aluminum, vial having an interior surface coated with a
30 polymer, in particular a fluorocarbon polymer. Advantageously other internal
surfaces, in particular such surfaces of components of the valve, or all of the
internal surfaces of the dispenser may be also coated with a polymer, in particular

a fluorocarbon polymer. Suitable fluorocarbon polymers include fluorocarbon polymers, which are made of multiples of one or more of the following monomeric units: tetrafluoroethylene (PTFE), fluorinated ethylene propylene (FEP), perfluoroalkoxyalkane (PFA), ethylene tetrafluoroethylene (ETFE), vinylidene fluoride (PVDF), and chlorinated ethylene tetrafluoroethylene. Polymers, which have a relatively high ratio of fluorine to carbon, such as perfluorocarbon polymers e.g. PTFE, PFA, and FEP, are preferred; FEP is particularly preferred.

The valve may be any suitable metering valve with an outlet made from, for example, stainless steel, acetal, nylon or polybutylene terephthalate and with seals made from nitrile or EPDM elastomer.

A formulation of the invention can be administered to the lung by oral or nasal inhalation. Oral inhalation is preferred, and conventional actuators for oral inhalation can be used in connection with a formulation of the invention. It has been found that good respirable doses are achieved with an orifice diameter within the range of 0.4 to 0.6 mm, preferably in the range 0.45 to 0.55 mm, most preferably 0.47 to 0.53 mm.

The invention will be illustrated by the following Examples.

Materials used:

α -lactose monohydrate supplied under the trade designation Pharmatose 325M by DMV International Pharma was micronised by fluid energy milling in a single pass (referred to here and in the following as "micronised lactose monohydrate"). Micronised lactose monohydrate (100 g) was dispersed in Anhydrous Ethanol (840 g) using a Silverson high shear mixer. This dispersion was added to the product reservoir of an Avestin Emulsiflex C50 homogenizer, and recirculated for 20 minutes at 10,000 psi. The dispersion was then passed out of the

homogenizer, after recirculation for 30 seconds, at 20,000 psi. Two separate batches were prepared and their particle size determined.

5 DL-alanine (10g, used as supplied by Fisher Chemicals, Loughborough UK) was dispersed in anhydrous ethanol (200g) using a Silversen high shear mixer set at 10,000 RPM for 1 minute. The resultant dispersion was poured into the product vessel of a M110EH Microfluidizer. The Microfluidiser was fitted with an auxiliary process module of 100 micron channel diameter and a G10Z diamond interaction chamber of 87 micron channel diameter. The G10Z chamber was fitted
0 downstream from the auxiliary process module. The dispersion was processed at 25,000 psi for 120 minutes.

Particle Size Analysis

5 For analysis of a Lactose/Ethanol slurry, a (0.5 ml) sample of the slurry, which was shaken for at least one minute to ensure homogeneity, was added to a solution of 0.05% Lecithin in Iso-octane (20 ml), and redispersed with mild ultrasonics for 1 minute.

20 The resulting suspension was introduced dropwise into the presentation cell (a Hydro 2000 SM small sample presentation cell) of a Malvern Mastersizer 2000TM laser diffraction particle sizer until the obscuration was in the working range (between 10 and 12 with a red laser), and left to circulate for 1 minute to allow complete mixing and thermal equilibrium to be established. Ten readings were
25 taken at 20 second intervals to establish that the particle size was stable. The General Purpose analysis model, as described in the Malvern Instruments Operators Guide, was used with refractive indices 1.533 (lactose), 1.392 (iso-octane) and absorbance 0.001 (lactose). The results are based on the average calculated results of 10 readings taken in succession. The procedure was
30 performed twice.

For analysis of Alanine, particle size analysis measurement was performed using a Malvern Mastersizer 2000™ laser diffraction particle sizer in an analogous manner to that described above for lactose, with the exception that a refractive index of 1.55 was employed for DL-alanine.

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Results of Particle Size Analysis by Malvern Mastersizer 2000

	Lactose	Lactose	DL-Alanine
	Example 1	Example 2	Example 3
Units	Microns	Microns	Microns
d(v,0.1)	0.075	0.071	0.077
d(v,0.5) median	0.181	0.160	0.190
d(v,0.9)	0.896	0.793	0.600
D[4,3] volume weighted mean	0.338	0.301	0.283
Units	Percent	Percent	Percent
vol under 0.05 micron	1.85	2.20	1.68
vol under 0.10 micron	20.78	24.35	19.13
vol under 0.20 micron	54.71	61.09	52.45
vol under 0.50 micron	81.33	82.53	86.56
vol under 1.0 micron	91.65	93.55	96.25
vol under 2.0 micron	99.38	99.84	99.98
vol under 5.0 micron	100.0	100.00	100.00
vol under 10.0 micron	100.00	100.00	100.00
vol under 20.0 micron	100.00	100.00	100.00

10 Exemplary Formulations

Examples 4 - 15

Each formulation was prepared in a PET vial. The appropriate amounts of micronised drugs, a racemic mixture of formoterol fumarate dihydrate and mometasone furoate, were weighed into the PET vial. A slurry of nanosized lactose in ethanol from Example 1 was added as a thick paste to the PET vial; but initially not allowed to mix with the powdered drugs. A propellant mixture was

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prepared in a separate aerosol can, containing the propellant and the balance of any other ingredients in proportion that had not already been added. A non-metering valve was crimped onto the PET vial, which was then charged with the prescribed weight of propellant mixture from the aerosol can to prepare
5 formulations by mixing all the ingredients. These were further mixed for 1 minute in an ultrasonic bath.

Comparative formulations A-D without lactose were prepared as follows. Each formulation was prepared in a PET vial. The appropriate amounts of
0 micronised drugs were weighed into the PET vial. A propellant mixture was prepared in a separate aerosol can, containing the propellant and the balance of any other ingredients in proportion that had not already been added (ie. a higher ethanol mix than that used for the examples of the invention). A non-metering valve was crimped onto the PET vial, which was then charged with the prescribed
5 weight of propellant mixture from the aerosol can to prepare formulations by mixing all the ingredients. These were further mixed for 1 minute in an ultrasonic bath.

The physical characteristics and suspension settling rates of prepared
20 formulations were examined by shaking the formulation and observing the formulation upon standing visually and by using an optical measuring technique such as that described in the Proceedings of Drug Delivery to the Lung VI p.10-13 (December 1995) printed by the Aerosol Society. To use such equipment, a PET vial containing a unit quantity of formulation was placed on a vortex mixer for 10
25 seconds and then immediately placed into the test apparatus. The apparatus was set up with an upper light beam and detector 3 mm below the liquid meniscus and a lower light beam and detector 3 mm above the base of the vial. It was also set up to automatically start recording the light transmission to each detector as the vial passed the upper beam upon insertion. The transmission was captured as a
30 plot of voltage against time. For sedimenting formulations the data from the upper sensor were used.

HFC 134a-based formulations

5	Formulation no.	A		4	
		<u>mg/ml</u>	<u>g/unit</u>	<u>mg/ml</u>	<u>g/unit</u>
	Formoterol fumarate	0.1320	0.0010	0.1320	0.0010
	Mometasone furoate	1.0000	0.0083	1.0000	0.0083
	Lactose monohydrate	0.0000	0.0000	1.3200	0.0109
	Oleic Acid	0.0606	0.0005	0.0606	0.0005
0	Ethanol	24.2200	0.2000	24.2200	0.2000
	HFA 134a	1185.5874	9.7902	1184.2674	9.7793

5	Formulation no.	5		6	
		<u>mg/ml</u>	<u>g/unit</u>	<u>mg/ml</u>	<u>g/unit</u>
	Formoterol fumarate	0.1320	0.0010	0.1320	0.0010
	Mometasone furoate	1.0000	0.0083	1.0000	0.0083
	Lactose monohydrate	1.9800	0.0164	2.6400	0.0218
	Oleic Acid	0.0606	0.0005	0.0606	0.0005
	Ethanol	24.2200	0.2000	24.2200	0.2000
20	HFA 134a	1183.6074	9.7738	1182.9474	9.7684

25	Formulation no.	B		7	
		<u>mg/ml</u>	<u>g/unit</u>	<u>mg/ml</u>	<u>g/unit</u>
	Formoterol fumarate	0.1320	0.0010	0.1320	0.0010
	Mometasone furoate	5.0000	0.0413	5.0000	0.0413
	Lactose monohydrate	0.0000	0.0000	1.3200	0.0109
	Oleic Acid	0.0606	0.0005	0.0606	0.0005
	Ethanol	24.2200	0.2000	24.2200	0.2000
30	HFA 134a	1181.5874	9.7572	1180.2674	9.7463

35	Formulation no.	8		9	
		<u>mg/ml</u>	<u>g/unit</u>	<u>mg/ml</u>	<u>g/unit</u>
	Formoterol fumarate	0.1320	0.0010	0.1320	0.0010
	Mometasone furoate	5.0000	0.0413	5.0000	0.0413
	Lactose monohydrate	1.9800	0.0164	2.6400	0.0218
	Oleic Acid	0.0606	0.0005	0.0606	0.0005
	Ethanol	24.2200	0.2000	24.2200	0.2000
	HFA 134a	1179.6074	9.7408	1178.9474	9.7354

The formulations containing nano-sized lactose show better physical stability over the comparative formulations. All the formulations containing nano-sized lactose exhibited a significant slower rate of sedimentation than the lactose-free
 5 formulations. After standing 15 minutes it was noted that the sediment volume of the nano-sized lactose formulations 4-6 occupied approximately 35 to 55% of the formulation volume, while the non-bulked formulations showed a sediment occupying about 20% of the total formulation volume.

0 A preparation of Formulation 6 was also carried out using the lab scale process described below for Formulation 16, and defined portions thereof were cold-transferred to the following container-closure systems:

- P Plain aluminium canister with a 50 mcl Spraymiser™ valve crimped.
 Q Aluminium canister lined with Fluorinated Ethylene Propylene polymeric
 5 coating, with a 50 mcl Spraymiser™ valve crimped.

Two of each of P and Q canisters were left to equilibrate at room temperature for 24 hours, then stored for 5 days at 50°C in a valve-down orientation, then allowed to equilibrate for 24 hours at room temperature prior to analysis. A High Pressure
 20 Liquid Chromatographic analysis of the canister contents was performed to assess the levels of impurities, with the following average results:

	P	Q
% impurities	1.26	0.98

25 The experiment showed that lining the internal surface of the canister with a fluorocarbon polymeric coating reduced the level of degradation significantly.

30 HFC 227-based formulations

Formulation no.	C	10
	<u>mg/ml</u>	<u>g/unit</u>
	<u>mg/ml</u>	<u>g/unit</u>

	Formoterol fumarate	0.1320	0.0009	0.1320	0.0009
	Mometasone furoate	1.0000	0.0072	1.0000	0.0072
	Lactose monohydrate	0.0000	0.0000	1.3200	0.0095
	Oleic Acid	0.0695	0.0005	0.0695	0.0005
5	Ethanol	27.8000	0.2000	27.8000	0.2000
	HFA 227	1360.9985	9.7914	1359.6785	9.7819

	Formulation no.	11		12	
		<u>mg/ml</u>	<u>g/unit</u>	<u>mg/ml</u>	<u>g/unit</u>
10	Formoterol fumarate	0.1320	0.0009	0.1320	0.0009
	Mometasone furoate	1.0000	0.0072	1.0000	0.0072
	Lactose monohydrate	1.9800	0.0143	2.6400	0.0190
	Oleic Acid	0.0695	0.0005	0.0695	0.0005
	Ethanol	27.8000	0.2000	27.8000	0.2000
15	HFA 227	1359.0185	9.7771	1358.3585	9.7724

	Formulation no.	D		13	
		<u>mg/ml</u>	<u>g/unit</u>	<u>mg/ml</u>	<u>g/unit</u>
	Formoterol fumarate	0.1320	0.0009	0.1320	0.0009
20	Mometasone furoate	5.0000	0.0360	5.0000	0.0360
	Lactose monohydrate	0.0000	0.0000	1.3200	0.0095
	Oleic Acid	0.0695	0.0005	0.0695	0.0005
	Ethanol	27.8000	0.2000	27.8000	0.2000
	HFA 227	1356.9985	9.7626	1355.6785	9.7531

	Formulation no.	14		15	
		<u>mg/ml</u>	<u>g/unit</u>	<u>mg/ml</u>	<u>g/unit</u>
	Formoterol fumarate	0.1320	0.0009	0.1320	0.0009
	Mometasone furoate	5.0000	0.0360	5.0000	0.0360
30	Lactose monohydrate	1.9800	0.0143	2.6400	0.0190
	Oleic Acid	0.0695	0.0005	0.0695	0.0005
	Ethanol	27.8000	0.2000	27.8000	0.2000
	HFA 227	1355.0185	9.7483	1354.3585	9.7436

35 The formulations without lactose exhibited a tendency to cream. The formulations containing nano-sized lactose exhibited superior physical stability properties, no creaming and a slow sedimentation rate. After standing 15 minutes it was noted that the sediment volume of the nano-sized lactose formulations occupied greater than 50% of the formulation volume, while the non-bulked formulations had

nebulous flocs in which more than half of the flocs creamed and others floated in the formulation without filling any particular region of the formulation volume. It was also observed that the sedimentation rate for the nano-bulked HFC 227 based formulations was slower than the corresponding formulations using HFC 134a.

These examples demonstrate that the use of a nano-sized bulking agent enhances the physical stability of dispersed formoterol and mometasone particles.

Exemplary formulation 16

A Lab scale preparation of each formulation was carried out to make 100 inhalers as follows. Oleic acid was weighed into a stainless steel beaker, followed by a quantity of the slurry of lactose with ethanol (from Example 2). Finally, the balance of ethanol was added and the mixture stirred with a spatula to ensure complete mixing. Each micronized drug, a racemic mixture of formoterol fumarate dihydrate and mometasone furoate, was weighed into another stainless steel beaker. The 2 beakers were transferred to the dehumidified atmosphere of a cold filler. About 250 grams of propellant were added to the excipients beaker to wash into the batching vessel. The remainder of the propellant required for the formulation was added to the batching vessel. The drugs were tipped from their beaker into the batching vessel, and a high shear mixer lowered in to mix all the ingredients at 8000 rpm for two minutes. A magnetic flea was added to the batching vessel to allow continued mixing while the batch was transferred to aerosol canisters.

Formulation no.	16	
	<u>mg/ml</u>	<u>g/unit</u>
Formoterol fumarate	0.1320	0.0010
Mometasone furoate	1.2500	0.0103
Lactose monohydrate	1.3200	0.0109
Oleic Acid	0.0606	0.0005

Ethanol	24.2200	0.2000
HFA 134a	1184.0174	9.7773

5 Formulation 16 was transferred into 10 ml aluminium canisters and crimped with 50 microlitre Spraymiser™ valves.

Test Methods

Uniformity of Content

10 The aerosol units to be tested were inserted sequentially into a single standard actuator. Each inhaler so assembled was shaken with a gentle rolling action through 180 degrees inversion at 1 cycle per second for 4 seconds. Within 2 seconds a single shot was fired to waste by pressing the unit down for 1 second, then the inhaler allowed to stand for 30 seconds. This process was repeated to
15 fire 10 "priming" shots to waste.

The apparatus described and illustrated in European Pharmacopoeia 3rd edition 2001 Supplement pages S1658-9 with a grommet to seal the actuator to the apparatus was used to collect shots fired for assay. Each inhaler was actuated
20 using the same actuation protocol as used for the priming shots. 5 seconds after actuation, the pump was switched off and the inhaler removed. The apparatus (with filter) and grommet were quantitatively rinsed with 10 ml of sample diluent (methanol : acetonitrile : water in the ratio 1:1:1) by sealing the 2 ends of the apparatus, shaking for 30 seconds, placing in an ultrasonic bath for 1 minute and
25 shaking for a further 30 seconds. The sample diluent was then analysed by High Pressure Liquid Chromatography to determine the content of both drugs. This method was used to assay the content of shots 11-16 (start), 59-62 (middle) and 118-120 (end) of each unit tested.

Loss of Dose

30 The loss of dose was determined by applying the method for content analysis above.

The protocol was as follows.

After collecting dose 16, each inhaler was left to stand for 24 hours in a valve-up orientation. Then without priming, shots 17 and 18 were analysed in the same way as shots 15 and 16. For the double shot values tabulated, 2 consecutive shots taken 24 hours after the previous dispensed dose were compared with shots 15 and 16, the values corresponding to the mean of shots 17 and 18 represented as a percentage of the mean of shots 15 and 16. The single shot values compared shot 17 with the mean of shots 15 and 16.

Andersen particle size

10 The Andersen sampler (Apparatus 1, US Pharmacopoeia 24 monograph <601>) was used as follows. A filter paper (Whatman 934-AH) was cut to fit the "F" stage, which was incorporated into the stack. The remainder of the stack was assembled, including the USP throat.

Each unit was tested as follows.

15 Priming shots (10) were fired through a standard actuator as described in the method for uniformity of content. The valve and valve stem interior were cleaned with water (30 ml) followed by ethanol (30 ml) and dried thoroughly. The unit was weighed. An unused sample actuator was used to dispense doses while the Andersen sampler was drawing air at 28.3 liter per minute. This was kept sealed into the port of the USP throat with a grommet until 5 seconds after actuation. 20 shots were fired, using the actuation protocol described for uniformity of content. The vacuum source was switched off 30 seconds after the final actuation. The unit was weighed.

25 The valve stem (including the interior), actuator, throat (including grommet), Stage 0 and inlet cone, plates 0-2, plates 3-5, plates 6-7 and the filter were separately rinsed with the following diluents:

Parts	Diluent
Valve stem	Acetonitrile (20 ml), methanol (20 ml), water (20 ml), sequentially
Actuator	
Throat and grommet	Acetonitrile (30 ml), methanol (30 ml), water (30 ml), sequentially

Stage 0 and inlet cone	Sample diluent (10 ml)
Plates 0-2	
Plates 3-5	Sample diluent (50 ml)
Plates 6-7	Sample diluent (10 ml)
Filter	Acetonitrile (3ml), methanol (3ml), water (3ml), sequentially

The sample diluent was as used for uniformity of content

Each washing was analysed by High Pressure Liquid Chromatography to determine each drug content in micrograms.

5 The calculation of parameters tabulated is as follows.

$$\text{Fine particle fraction} = \frac{\text{drug analysed from plates 3 to 7 and the filter (micrograms)}}{\text{Drug analysed from the throat, grommet, Stage 0, inlet cone, Plates 0 to 7 and the filter (micrograms)}} \times 100\%$$

10

$$\text{Fine particle dose} = \text{drug analysed from plates 3 to 7 and the filter (micrograms)}$$

$$\text{Mean metered dose} = \text{total recovery of drug from all diluent samples (micrograms)}$$

Results

15

Uniformity of Content (start, middle and end)

Dose dispensed per shot ex actuator in micrograms and relative standard deviation expressed as %

Micrograms per actuation		Start	Middle	End
Formoterol fumarate	mcg/ actuation	5.5	6.2	5.8
	RSD%	5.4	6.0	5.0
Mometasone furoate	mcg/ actuation	53.2	55.4	54.5
	RSD%	6.1	5.4	3.7

20

The uniformity of content is satisfactory.

Loss of Dose test (% dispensed after 24 hours standing compared with 0 hours)

%		
Formoterol fumarate	Single shot	101.5
	Double shot	106.0
Mometasone furoate	Single shot	95.5
	Double shot	99.1

The formulation has performed well after standing for a 24 hour period.

5

10

Andersen particle size

	Formoterol fumarate	Mometasone furoate
Mean fine particle fraction (%)	39.4	40.7
Fine particle dose (micrograms per shot)	2.3	21.6
Mean metered shot (micrograms)	7.3	66.4

15 Formulations 17 and 18

These formulations were prepared as described under the heading Formulation 16.

Formulation no.	E		17		18	
	mg/ml	g/unit	mg/ml	g/unit	mg/ml	g/unit
Formoterol fumarate	0.1320	0.0009	0.1320	0.0009	0.1320	0.0009
Mometasone furoate	1.2500	0.0090	1.2500	0.0090	5.0000	0.0360
Lactose monohydrate	0.0000	0.0000	1.3200	0.0095	1.3200	0.0095
Oleic Acid	0.0695	0.0005	0.0695	0.0005	0.0695	0.0005

20

Ethanol	27.8000	0.2000	27.8000	0.2000	27.8000	0.2000
HFA 227	1359.4285	9.7801	1360.7485	9.7896	1355.6785	9.7531

- 5 Formulations 17, 18 and E were transferred into 10 ml FEP-lined aluminium canisters and crimped with 50 microlitre valves incorporating a thermoplastic elastomer gasket to prevent contact between the formulation and aluminium surfaces, as described and shown in fig.1 of US 6264923 B1.

10

15 Results

Loss of Dose test (% dispensed after 24 hours standing compared with 0 hours)

%	Formulation	E	17	18
Formoterol fumarate	Single shot	82.2	101.1	94.6
	Double shot	96.1	106.8	97.2
Mometasone furoate	Single shot	84.0	99.5	98.9
	Double shot	98.1	104.0	100.9

- 20 The loss of dose for formulations 17 and 18 was far superior to that for the comparative formulation E.

Andersen particle size

Formulation	17		18	
	Formoterol fumarate	Mometasone furoate	Formoterol fumarate	Mometasone furoate
Mean fine particle fraction (%)	47.1	47.6	40.2	43.0

Fine particle dose (micrograms per shot)	2.6	24.6	2.2	87.6
Mean metered shot (micrograms)	6.8	62.5	6.6	248.5

The Andersen particle size results represent reasonable delivery from the formulations.

5

Examples 19 and 20

A formulation according to Formulation 16 was prepared and defined portions were transferred to PET vials, which were subsequently crimped with a non-metering valve, in a similar manner as that described for Examples 4-15. This formulation is designated as Formulation 19. Formulation 20 was prepared in the same way as formulation 16, but with Alanine (from Example 3) substituted for Lactose. A comparative formulation, Formulation F was prepared similarly to formulation E. Corresponding portions of these formulations were also transferred to PET vials. The prepared formulations were then examined visually and by using an optical measuring technique as described above for Examples 4-15.

Formulation no.	F		19		20	
	mg/ml	g/unit	mg/ml	g/unit	mg/ml	g/unit
Formoterol fumarate	0.1320	0.0010	0.1320	0.0010	0.1320	0.0010
Mometasone furoate	1.2500	0.0103	1.2500	0.0103	1.2500	0.0103
Lactose monohydrate	0.0000	0.0000	1.3200	0.0109	0.0000	0.0000
Alanine	0.0000	0.0000	0.0000	0.0000	1.2300	0.0101
Oleic Acid	0.0606	0.0005	0.0606	0.0005	0.0606	0.0005
Ethanol	24.2200	0.2000	24.2200	0.2000	24.2200	0.2000
HFA 134a	1185.3374	9.7882	1184.0174	9.7781	1184.0174	9.7781

Formulation 19 showed a significantly slower settling rate than comparative formulation F, while surprisingly Formulation 20 sedimented even more slowly than Formulation 19. After standing 15 minutes it was noted that the sediment volume of the nano-sized lactose and alanine formulations occupied

approximately 30% and 23% respectively of the formulation volume, while the non-bulked formulation showed a sediment occupying about 7% of the total formulation volume.

CLAIMS

1. A pharmaceutical aerosol formulation comprising particles of (a) formoterol or a pharmaceutically acceptable salt, solvate or physiologically functional derivative thereof and (b) mometasone or a pharmaceutically acceptable salt,
5 solvate, or physiologically functional derivative thereof dispersed in a propellant selected from the group consisting of 1,1,1,2-tetrafluoroethane, 1,1,1,2,3,3,3-heptafluoropropane and a mixture thereof, and a bulking agent having a mass median diameter of less than one micron.
2. A pharmaceutical aerosol formulation according to claim 1, wherein the
10 formoterol is in the form of formoterol fumarate.
3. A pharmaceutical aerosol formulation according to claim 2, wherein the formoterol is in the form of formoterol fumarate dihydrate.
4. A pharmaceutical aerosol formulation any one of claims 1 to 3, wherein the mometasone is in the form of mometasone furoate.
- 15 5. A pharmaceutical aerosol formulation according to any one of claims 1 to 4, wherein the formoterol is present in an amount of about 0.06 to 0.60 mg per ml.
6. A pharmaceutical aerosol formulation according to any one of claims 1 to 5, wherein the mometasone is present in amount of about 0.5 to 15.0 mg per ml.
- 20 7. A pharmaceutical aerosol formulation according to any one of claims 1 to 6, wherein the bulking agent is selected from groups consisting of ascorbic acid, saccharides, polysaccharides, amino acids, organic and inorganic salts, urea and propylidone.
8. A pharmaceutical aerosol formulation according to claim 7, wherein the
25 bulking agent is selected from lactose, DL-alanine, glucose, D-galactose, D(+)trehalose dihydrate, sucrose, maltose, D(+)raffinose pentahydrate, sodium saccharin, starches, modified celluloses, dextrans, glycine, sodium chloride, calcium carbonate, sodium tartrate and calcium lactate.

9. A pharmaceutical aerosol formulation according to claim 7 or claim 8, wherein the bulking agent is lactose.
10. A pharmaceutical aerosol formulation according to any one of claims 1 to 9, wherein the weight ratio of formoterol to bulking agent is in the range 1:0.1 to 1:30.
11. A pharmaceutical aerosol formulation according to any one of claims 1 to 10, wherein the bulking agent has a mass median diameter of not more than 300 nm.
12. A pharmaceutical aerosol formulation according to any one of claims 1 to 11, wherein the formulation further comprises a surfactant.
13. A pharmaceutical aerosol formulation according to any one of claims 1 to 12, wherein the formulation further comprises ethanol.
14. A pharmaceutical aerosol formulation according to claim 13, wherein ethanol is present in amount of from 0.1 to 5% by weight of the formulation.
15. A dispenser comprising an aerosol vial equipped with a dispensing valve, said aerosol vial containing a formulation according to any one of claims 1 to 14.
16. A dispenser according to claim 16, wherein an interior surface of the aerosol vial is coated with a fluorocarbon polymer.
17. A method of preparing a formulation according to claim 1, the method comprising the steps of (i) forming a slurry of bulking agent with a component of the formulation; (ii) subjecting the slurry to high pressure homogenization; and (iii) combining the resulting slurry with other components of the aerosol formulation.

INTERNATIONAL SEARCH REPORT

Intern Application No

PCT/US 03/08710

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K9/12 A61K9/70 A61K31/58 A61K31/167 A61K47/10
 A61K47/26 A61P11/06 A61K47/18

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, CHEM ABS Data, MEDLINE, EMBASE, BIOSIS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	<p>WO 03 020253 A (HART JOHN L ; SCHERING CORP (US); SEQUEIRA JOEL A (US); SHARPE STEF) 13 March 2003 (2003-03-13) the whole document examples 7-9 page 14, paragraph 1 claims 10-19</p> <p style="text-align: center;">--- -/-</p>	1-17



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

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Date of the actual completion of the international search

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Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

Luangkhot, N

INTERNATIONAL SEARCH REPORT

Intern. Application No.

PCT/US 03/08710

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 02 30394 A (JINKS PHILIP A ;LISTER JAMES T (GB); MCKENZIE LESLEY (GB); 3M INNO) 18 April 2002 (2002-04-18) cited in the application the whole document claims 1-10 claims page 4, line 3-24 page 5, line 7-22 page 7, line 10-17 page 7, line 19 -page 8, line 22 page 10, line 13-18 page 11, line 11-17 page 13, line 14-16 page 16, line 14-17 examples ----	1-17
Y	WO 00 51591 A (NOVARTIS ERFIND VERWALT GMBH ;NOVARTIS AG (CH); CLARKE JEREMY GUY) 8 September 2000 (2000-09-08) cited in the application the whole document page 3, paragraph 5 -page 4, paragraph 1 ----	1-17
X	WO 97 47286 A (JINKS PHILIP A ;OLIVER MARTIN J (GB); PALING SIMON G (GB); JAISWAL) 18 December 1997 (1997-12-18) the whole document page 4, line 12 -page 6, line 25; examples 16-33 page 11, line 14-17 page 12, line 20 -page 13, line 3 claims 11-23 ----	1-17
Y		1-17
A	WO 00 07567 A (HERZOG KURT ;JAGO RES AG (CH); KRAUS HOLGER (CH); MUELLER WALZ RUD) 17 February 2000 (2000-02-17) the whole document page 12, paragraph 2 page 19, paragraph 2 -----	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

Internal Application No

PCT/US 03/08710

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 03020253 A	13-03-2003	WO 03020253 A2 US 2003114428 A1	13-03-2003 19-06-2003
WO 0230394 A	18-04-2002	AU 1131102 A CA 2425035 A1 EP 1324749 A2 NO 20031597 A WO 0230394 A2	22-04-2002 18-04-2002 09-07-2003 30-05-2003 18-04-2002
WO 0051591 A	08-09-2000	AT 241970 T AU 3284300 A BR 0008699 A CA 2362499 A1 CN 1341017 T CZ 20013168 A3 DE 60003160 D1 WO 0051591 A1 EP 1156790 A1 EP 1325765 A1 HU 0200142 A2 JP 2002538110 A NO 20013988 A NZ 513742 A SK 12302001 A3 US 2003050290 A1 US 2002055494 A1	15-06-2003 21-09-2000 26-12-2001 08-09-2000 20-03-2002 12-12-2001 10-07-2003 08-09-2000 28-11-2001 09-07-2003 29-06-2002 12-11-2002 16-08-2001 28-09-2001 03-12-2001 13-03-2003 09-05-2002
WO 9747286 A	18-12-1997	AU 726382 B2 AU 3373997 A CA 2257841 A1 EP 0934057 A1 JP 2000513340 T NO 985720 A NZ 333202 A US 6054488 A WO 9747286 A1 ZA 9704546 A	02-11-2000 07-01-1998 18-12-1997 11-08-1999 10-10-2000 11-02-1999 23-06-2000 25-04-2000 18-12-1997 23-11-1998
WO 0007567 A	17-02-2000	AT 234604 T AU 749697 B2 AU 4893999 A CA 2338680 A1 WO 0007567 A1 CN 1315852 T DE 59904648 D1 DK 1102579 T3 EP 1102579 A1 JP 2002522374 T NO 20010531 A NZ 509489 A US 6475467 B1	15-04-2003 04-07-2002 28-02-2000 17-02-2000 17-02-2000 03-10-2001 24-04-2003 14-07-2003 30-05-2001 23-07-2002 31-01-2001 25-10-2002 05-11-2002